IN THE UNITED STATES DISTRICT COURT FOR THE DISTRICT OF KANSAS

BARBARA BAUGHN,)	
)	
	Plaintiff,)	
)	CIVIL ACTION
v.)	
)	No. 03-2626-KHV
ELI LILLY AND COMPANY,)	
)	
	Defendant.)	
)	

ORDER

Barbara Baughn filed this product liability action against Eli Lilly and Company ("Lilly"). Plaintiff alleges that she suffered injuries because her mother took diethylstilbestrol ("DES"), a prescription drug, during her pregnancy with Barbara in 1964 and 1965. This matter is before the Court on Defendant Eli Lilly And Company's Second Motion For Summary Judgment (Doc. #42) filed October 1, 2004; Plaintiff's Motion To Strike Affidavit Of John F. Kuckelman And Supporting Memorandum Of Law (Doc. #44) filed October 11, 2004; Plaintiff's Motion To Strike Second Affidavit Of John F. Kuckelman And Supporting Memorandum Of Law (Doc. #61) filed December 10, 2004; and Plaintiff's Motion To Limit Or Exclude Defendant's Expert Testimony And Memorandum Of Law In Support Thereof (Doc. #41) filed October 1, 2004. For reasons stated below, the Court overrules all of the motions.

Summary Judgment Standards

Summary judgment is appropriate if the pleadings, depositions, answers to interrogatories, and admissions on file, together with the affidavits, if any, show no genuine issue as to any material fact and that the moving party is entitled to a judgment as a matter of law. See Fed. R. Civ. P. 56(c); accord Anderson

v. Liberty Lobby, Inc., 477 U.S. 242, 247 (1986); Vitkus v. Beatrice Co., 11 F.3d 1535, 1538-39 (10th Cir. 1993). A factual dispute is "material" only if it "might affect the outcome of the suit under the governing law." Anderson, 477 U.S. at 248. A "genuine" factual dispute requires more than a mere scintilla of evidence. Id. at 252.

The moving party bears the initial burden of showing the absence of any genuine issue of material fact. Celotex Corp. v. Catrett, 477 U.S. 317, 323 (1986); Hicks v. City of Watonga, 942 F.2d 737, 743 (10th Cir. 1991). Once the moving party meets its burden, the burden shifts to the nonmoving party to demonstrate that genuine issues remain for trial "as to those dispositive matters for which it carries the burden of proof." Applied Genetics Int'l, Inc. v. First Affiliated Sec., Inc., 912 F.2d 1238, 1241 (10th Cir. 1990); see also Matsushita Elec. Indus. Co. v. Zenith Radio Corp., 475 U.S. 574, 586-87 (1986); Bacchus Indus., Inc. v. Arvin Indus., Inc., 939 F.2d 887, 891 (10th Cir. 1991). The nonmoving party may not rest on its pleadings but must set forth specific facts. Applied Genetics, 912 F.2d at 1241.

"[W]e must view the record in a light most favorable to the parties opposing the motion for summary judgment." <u>Deepwater Invs.</u>, Ltd. v. Jackson Hole Ski Corp., 938 F.2d 1105, 1110 (10th Cir. 1991). Summary judgment may be granted if the non-moving party's evidence is merely colorable or is not significantly probative. <u>Anderson</u>, 477 U.S. at 250-51. "In a response to a motion for summary judgment, a party cannot rely on ignorance of facts, on speculation, or on suspicion, and may not escape summary judgment in the mere hope that something will turn up at trial." <u>Conaway v. Smith</u>, 853 F.2d 789, 794 (10th Cir. 1988). Essentially, the inquiry is "whether the evidence presents a sufficient disagreement to require submission to the jury or whether it is so one-sided that one party must prevail as a matter of law." <u>Anderson</u>, 477 U.S. at 251-52.

Factual Background

The following facts are uncontroverted, deemed admitted or, where disputed, viewed in the light most favorable to plaintiffs.

Barbara Baughn was born on August 19, 1965 in Chanute, Kansas, and has lived in Kansas her entire life. Plaintiff's mother, who was pregnant with Barbara in 1964 and 1965, testified that plaintiff was born one week before her due date. From the second month of pregnancy until plaintiff was born, plaintiff's mother took DES prescribed by Dr. Reuben Burkman. Plaintiff's mother purchased DES at Baker & Burkman Pharmacy in Chanute, Kansas. Dr. Burkman owned Baker & Burkman Pharmacy and the pharmacy was attached to his clinic. Plaintiff's mother took DES four times a day in small round red tablets.

In 1964 and 1965, Lilly made and sold DES in a 25 milligram dosage in round red enteric coated tablets. Lilly's product information recommended a dosage of 25 to 100 milligrams per day for prevention of "accidents of pregnancy."

In 1965, Dr. Burkman prescribed DES on a fairly common basis to prevent miscarriage. Joan Augustine, a nurse, began working for Dr. Burkman in the spring of 1965. Augustine testified that on occasion, she went into the pharmacy to "fix up" a prescription when the pharmacist was gone. Augustine

Lilly has submitted various documents which purport to show that in 1965, more than 95 companies manufactured DES. Plaintiff seeks to exclude these documents as hearsay. See Plaintiff's Motion To Strike Affidavit Of John F. Kuckelman And Supporting Memorandum Of Law (Doc. #44) filed October 11, 2004. For purposes of defendant's summary judgment motion, however, the Court need not consider whether other companies manufactured DES. The relevant issue is whether any manufacturer beside Lilly supplied DES to the Baker & Burkman Pharmacy. Accordingly, the Court overrules as moot plaintiff's motion to strike (Doc. #44).

does not recall when she began helping with prescriptions, but she probably did so for some prescriptions in 1965. Augustine cannot recall preparing a prescription for plaintiff's mother – or any other DES prescription – in 1965.

Augustine testified that the pharmacy stocked Lilly DES in little red enteric-coated tablets in 1965, that she recalls only DES from Lilly on the pharmacy shelf, that she believes that the pharmacy carried only DES from Lilly, and that the pharmacy carried a lot of Lilly products. Augustine testified that she cannot recall specifically what dosage the Lilly red DES tablet came in, but that she believed it was one milligram.

Dr. Burkman always had a Physicians' Desk Reference ("PDR") in each examining room. In the PDRs for 1964 and 1965, Lilly's DES advertisements do not mention any risk related to DES during pregnancy, either to the mother or the fetus.²

Plaintiff claims that as a result of her exposure to DES *in utero*, she suffered pregnancy complications, pregnancy losses and infertility for which she sought medical treatment in Kansas. Barbara did not learn of her claim against defendant until late 2001. On May 8, 2003, Barbara Baughn filed suit against Lilly in the United States District Court for the District of Columbia. On November 7, 2003, pursuant to 28 U.S.C. § 1404(a), the court transferred its case to this Court.

Plaintiff asserts four theories of recovery in this product liability action: negligence, strict liability,

In the 1964 and 1965 editions of the PDR, the brand name index lists only Lilly as a manufacturer of "Diethylstilbestrol." Lilly has submitted various documents which purport to show that in the 1964 and 1965 PDRs, other companies are listed as manufacturers of "DES" and that in 1964 and 1965 other companies manufactured DES in red enteric coated tablets in a 25 milligram dosage. Plaintiff seeks to exclude these documents as hearsay. Plaintiff's Motion To Strike Second Affidavit Of John F. Kuckelman And Supporting Memorandum Of Law (Doc. #61) filed December 10, 2004. As explained above, for purposes of this motion, the Court need not consider whether other companies manufactured DES. Accordingly, the Court overrules as moot plaintiff's motion to strike (Doc. #61).

breach of warranty and negligent misrepresentation. The parties and the Court agree that Kansas substantive law applies in this case.

In its motion for summary judgment, Lilly argues that (1) plaintiff cannot prove that it manufactured the DES to which she was allegedly exposed and (2) absent proof that Dr. Burkman prescribed only DES manufactured by Lilly, plaintiff cannot rely on the presumption that he would have heeded a different warning by Lilly.

Analysis

I. Lilly As Manufacturer Of The DES Which Plaintiff's Mother Ingested

Defendant argues that plaintiff cannot prove that it manufactured the DES to which she was allegedly exposed. The Court disagrees. Augustine testified that Dr. Burkman commonly prescribed DES in 1965, that the pharmacy stocked Lilly DES in 1965, and that the pharmacy carried a lot of Lilly products. See Augustine Depo. at 41-42, 50-56. Lilly argues that plaintiff cannot establish that in 1965, the pharmacy carried Lilly DES in a 25 milligram tablet. Augustine testified that she cannot recall specifically what dosage the Lilly red DES tablet came in, but that she believes it was one milligram. See Augustine Depo. at 52. Augustine did not exclude the possibility that Lilly's DES was available at the pharmacy in other dosages. Despite Augustine's uncertainty about the precise dosages of DES which the pharmacy dispensed, she recalls only Lilly DES on the pharmacy shelf and she believes that the pharmacy only carried DES from Lilly. Augustine Depo. at 50-56. Combined with plaintiff's mother's testimony that she took a red DES tablet in a 25 milligram dosage which she obtained from the Baker & Burkman Pharmacy, a reasonable jury could conclude that Lilly manufactured the DES which plaintiff's mother ingested in 1965. The Court therefore overrules this portion of defendant's motion for summary judgment.

II. Proximate Cause

In a closely related argument, Lilly argues that absent proof that Dr. Burkman prescribed only Lilly's DES, plaintiff cannot rely on the presumption that Dr. Burkman would have heeded a different warning by Lilly. In order to survive summary judgment, plaintiff must provide evidence that a failure to warn proximately caused her injuries. See Wooderson v. Ortho Pharm. Corp., 235 Kan. 387, 409, 681 P.2d 1038, 1057 (1984); see also Eck v. Parke, Davis & Co., 256 F.3d 1013, 1018 (10th Cir. 2001). Kansas law allows a rebuttable presumption of causation once plaintiff establishes that a warning is inadequate. If plaintiff proves that Lilly failed to provide a proper warning, Kansas law presumes that a doctor using that product would have heeded a proper warning. See Wooderson, 235 Kan. at 407, 681 P.2d at 1057. Essentially, the law presumes that but for the inadequate warning, the patient would not have been harmed, since the doctor would have given the patient an adequate warning if the doctor had ever received it, and that the inadequate warning is therefore the cause of the patient's injury. See id. at 409, 681 P.2d at 1057. Defendant may rebut this presumption by establishing that although the prescribing physician would have read and heeded the warning or additional information, the warning would not have changed the course of treatment. See Eck, 256 F.3d at 1019. If Lilly provides credible evidence to rebut the presumption, the presumption disappears and the burden shifts back to plaintiff to affirmatively prove causation. See id.; Woulfe v. Eli Lilly & Co., 965 F. Supp. 1478, 1483 (E.D. Okla. 1997).

For purposes of this motion, Lilly does not dispute that its warning was inadequate. Lilly argues, however, that plaintiff is not entitled to a heeding presumption because she cannot establish that in 1965, Dr. Burkman prescribed only DES manufactured by Lilly. As explained above, plaintiff has presented evidence which raises a genuine issue of material fact whether Lilly manufactured the DES which her

mother ingested in 1965. In particular, Augustine testified that she recalls only Lilly's DES on the pharmacy shelf and that she believed that the pharmacy only carried DES from Lilly. See Augustine Depo. at 50-56. Based on such testimony, a reasonable jury could also conclude that in 1965, Dr. Burkman prescribed DES manufactured only by Lilly. The Court therefore overrules this portion of defendant's motion for summary judgment.

III. Claim On Behalf Of Derek Baughn

Kansas law permits individuals to assert loss of consortium claims on behalf of their spouses.

K.S.A. § 23-205 provides in part as follows:

Where, through the wrong of another, a married person shall sustain personal injuries causing the loss or impairment of his or her ability to perform services, the right of action to recover damages for such loss or impairment shall vest solely in such person, and any recovery therefor, so far as it is based upon the loss or impairment of his or her ability to perform services in the household and in the discharge of his or her domestic duties, shall be for the benefit of such person's spouse so far as he or she shall be entitled thereto.

K.S.A. § 23-205. Defendant argues that plaintiff cannot assert a loss of consortium claim because at the time of her injuries, she was not married. Defendant relies solely on <u>Dixon v. CertainTeed Corp.</u>, 915 F. Supp. 1158 (D. Kan. 1996). <u>Dixon</u> noted that for a plaintiff to recover for loss of consortium, "he first must demonstrate the existence of a valid marriage contract at the time of his accident pursuant to K.S.A. § 23-205." <u>Id.</u> at 1159. <u>Dixon</u> involved a slip and fall at defendant's plant where plaintiff asserted that he satisfied K.S.A. § 23-205 through a common-law marriage. <u>Dixon</u> did not address when an injury must be suffered by a spouse to permit a claim under K.S.A. § 23-205, and plaintiff in <u>Dixon</u> did not assert latent injuries first discovered after the marriage.

Neither party has briefed whether the statutory language of K.S.A. § 23-205 permits a loss of

consortium claim for latent injuries discovered after an individual is married but caused by defendant's acts before the individual married. Based solely on the language of K.S.A. § 23-205, the Court cannot conclude that such a claim is precluded. Several courts have permitted loss of consortium claims in the latent injury context. See Green v. A.P.C. (Am. Pharm. Co.), 960 P.2d 912, 918-19 (Wash. 1998) (loss of consortium claim based on wife's exposure in utero to DES; injuries did not manifest themselves until after marriage); Aldredge v. Whitney, 591 So.2d 1201, 1205 (La. Ct. App. 1991) (loss of consortium claim based on back injuries caused by pre-marriage tortious act that manifested themselves during marriage); Kociemba v. G.D. Searle & Co., 683 F. Supp. 1577, 1578 (D. Minn. 1988) (loss of consortium claim based on wife's infertility caused by IUD implanted before marriage; infertility did not manifest itself until after marriage); Furby v. Raymark Indus., Inc., 397 N.W.2d 303, 306-07 (Mich. Ct. App. 1986) (loss of consortium claim based on asbestos exposure before marriage); Stager v. Schneider, 494 A.2d 1307, 1316 (D.C. 1985) (loss of consortium claim based on radiologist's negligence before marriage). But see Zwicker v. Altamont Emergency Room Physicians Med. Group, 118 Cal. Rptr.2d912 (Cal. Ct. App. 2002); Fullerton v. Hosp. Corp. of Am., 660 So.2d 389, 390-91 (Fla. Dist. Ct. App. 1995); Anderson v. Eli Lilly & Co., 588 N.E.2d 66, 68 (N.Y. 1991). Defendant does not explain why the Court should bar plaintiff's loss of consortium claim where, at the time of marriage, plaintiff's own claim had not accrued because she had not suffered ascertainable injuries. Moreover, although plaintiff apparently suffered "substantial injury" in 1965 when she was exposed to DES, she may be able to show that she suffered additional injuries when she attempted to become pregnant after she married. For these reasons, the Court overrules defendant's motion for summary judgment on plaintiff's claim under K.S.A. § 23-205.

IV. Consolidated Product Liability Claim

Plaintiff asserts four theories of recovery: negligence, strict liability, breach of warranty and negligent misrepresentation. The underlying purpose of the Kansas Product Liability Act ("KPLA"), K.S.A § 60-3301 et seq., is "to consolidate all product liability actions, regardless of theory, into one theory of legal liability." Patton v. Hutchinson Wil-Rich Mfg. Co., 253 Kan. 741, 756, 861 P.2d 1299, 1311 (1993); see Savina v. Sterling Drug, Inc., 247 Kan. 105, 126, 795 P.2d 915, 931 (1990) (KPLA applies to actions based on strict liability in tort as well as negligence, breach of express or implied warranty, and breach of or failure to discharge a duty to warn or instruct); Fennesy v. LBI Mgmt., Inc., 18 Kan. App.2d 61, 66, 847 P.2d 1350, 1355 (1993) (purpose of KPLA to merge all legal theories of product liability into single product liability claim). Kansas law, however, recognizes that a product can be defective in one of three ways: (1) a manufacturing defect, i.e. a flaw in the manufacturing of the product; (2) a warning defect, i.e. a failure to adequately warn of a risk or hazard related to the product design; or (3) a design defect, i.e. a product which although perfectly manufactured contains a defect that makes it unsafe. Savina v. Sterling Drug, Inc., 247 Kan. 105, 114, 795 P.2d 915, 923 (1990); Hiner v. Deere & Co., 161 F. Supp.2d 1279, 1282-83 (D. Kan. 2001), rev'd in part on other grounds, 340 F.3d 1190 (10th Cir. 2003).

Each of plaintiff's four theories appears to rely primarily on a warning defect. Under the KPLA, plaintiff can only assert a single product liability claim based on that theory. Liberally construing the pretrial order, however, plaintiff also asserts negligence and strict liability claims based on manufacturing and design defects. Neither party has addressed the issue, but it appears that plaintiff is asserting three separate theories of product liability: manufacturing, warning and design defects. Defendant correctly notes that all

theories of recovery (negligence, strict liability, breach of warranty and negligent misrepresentation) merge into one legal theory called a "product liability claim," but defendant ignores the fact that plaintiff may allege multiple defects in a single product. The pretrial order does not precisely assert a single claim based on multiple defects, but the Court finds that it is unnecessary to modify the pretrial order at this stage. The parties can submit proposed jury instructions which reflect the legal limitation on multiple claims based on the same product defect. Accordingly, the Court overrules defendant's motion for summary judgment on this issue.

V. Plaintiff's Motion To Exclude Expert Testimony

Plaintiff seeks to exclude the deposition testimony of three defense experts who are deceased, Don Carlos Hines, M.D., Theodore G. Klumpp, M.D., and Edith L. Potter, M.D. The deposition testimony of these experts was taken in connection with various lawsuits in the late 1970s and early 1980s. Plaintiff argues that the deposition testimony likely will not satisfy the standards under Rules 702 through 705, Daubert v. Merrell Dow Pharms., Inc., 509 U.S. 579 (1993), and Kumho Tire Co. v. Carmichael, 526 U.S. 137 (1999). Plaintiff maintains that because defendant did not provide transcripts of the depositions before the deadline to file her Daubert motion, she could not more fully challenge the expert opinions. Plaintiff seeks leave to supplement her motion after defendant provides copies of the expert depositions.

Plaintiff apparently did not seek copies of the deposition transcripts until shortly before the motion to exclude expert testimony was due. Moreover, until the day that the <u>Daubert</u> motion was due, plaintiff did not notify defense counsel that the transcripts were necessary to prepare that motion. Defendant provided copies of the expert depositions on October 4, 2004 (the business day after plaintiff filed her motion to exclude). Plaintiff's reply brief on her motion to exclude was due October 26, 2004, <u>see</u> D.

Kan. Rule 6.1(d)(1), but she did not file a reply. The Court therefore has no record of the relevant deposition testimony, and plaintiff has not advanced specific grounds for excluding the expert testimony of Drs. Hines, Klumpp and Potter. Based solely on plaintiff's conclusory motion and absent copies of the deposition transcripts, the Court must overrule plaintiff's motion.³

IT IS THEREFOREORDERED that <u>Defendant Eli Lilly And Company's Second Motion For Summary Judgment</u> (Doc. #42) filed October 1, 2004 be and hereby is **OVERRULED**.

IT IS FURTHER ORDERED that Plaintiff's Motion To Strike Affidavit Of John F. Kuckelman

And Supporting Memorandum Of Law (Doc. #44) filed October 11, 2004 be and hereby is

OVERRULED.

IT IS FURTHER ORDERED that Plaintiff's Motion To Strike Second Affidavit Of John F.

Kuckelman And Supporting Memorandum Of Law (Doc. #61) filed December 10, 2004 be and hereby is OVERRULED.

IT IS FURTHER ORDERED that <u>Plaintiff's Motion To Limit Or Exclude Defendant's Expert</u>

<u>Testimony And Memorandum Of Law In Support Thereof</u> (Doc. #41) filed October 1, 2004 be and hereby is **OVERRULED**.

Because plaintiff did not file a reply some 22 days after receiving copies of the deposition transcripts, the Court overrules plaintiff's request to supplement her motion to exclude based on the deposition transcripts.

IT IS FURTHER ORDERED that trial is set for July 5, 2005 at 9:30 a.m.

Dated this 17th day of February, 2005 at Kansas City, Kansas.

s/ Kathryn H. Vratil
KATHRYN H. VRATIL
United States District Judge